Amendments to the Claims:

This listing will replace all prior versions and listings of claims in the application:

Listing of Claims:

1. (previously presented) A compound according to formula I:

wherein:

 R^1 and R^3 are each independently H, lower alkyl, -SO₃H, or -PO₃H₂, ; R^2 is selected from hydrogen, -SO₃H or -PO₃H; and X^1 is bound in the 2- or 3- position and is of the formula :

Ar-X3 T wherein -

Ar is furanyl, thienyl, pyridyl, eyelohexyl or benzyl and X3 is H. C. N. NR'. NR'R", NR'SO2 R", or O; wherein R' and R" are each independently H, or lower alkyl; and OR¹ is O(CH₂)_nY, wherein n is 1 to 2, Y is OR⁴, NR⁵R⁶, COOR⁴, or CONR⁵R⁶: or O[CH2CH (OH) CH2]Y, wherein Y is H, OR4, NR5R6, COOR4, or CONR5R6; wherein T is Y or [CH2CH (OH) CH2]Y, Y is H, OR⁴, NR⁵R⁶, COOR⁴, or CONR⁵R⁶ wherein R⁴, R⁵, and R⁶ are each independently H, or lower alkyl, and R⁵ and R⁶ together may form a 5 to 7membered ring; or pharmaceutically acceptable salts thereof, or Ar is phenyl and X³ is selected from the group consisting of NR'R", NHR', hydroxyl. Alkoxy, carbooxyalkoxyalkoxy, hydroxyl(hydroxyalkyl)alkoxy and N,N-dialkylaminoalkoxy, wherein R' and R' are each independently selected from hydrogen, alkyl, N,N-(dialkylamino)alkyl, alkoxyalkyl, hydroxyalkyl, alkylsulfonyl and alkylsulfonato alkyl asubstituent on the ortho, meta or para position of the phenyl ring and is C. N. NR'SO2R" or O: wherein R' and R'' are each independently H or lower alkyl; and OR is O(CH2)nY wherein n is 1 or 2 Y is OR4, NR5R6, COOR4 or CONR5R6 or O/CH_CH_(OH)CH_1Y wherein Y is H_OR4_NR5R6_COOR4 or CONR5R6 wherein T is Y or OICH-CH-(OH)CH-IV. Y-is-H. OR4. NR3R6. COOR4 or CONR5R6 wherein R4. R5 and R6are each independently H or lower alkyl and R⁵ and R⁶ together may form a 5 to 7 membered ring or pharmaceutically acceptable salts thereof, subject to the provise that X3T is not OR'

or NR'R" wherein R' and R" arre each independently H or lower alkyl.

- 2. (Cancelled)
- 3. (Cancelled)
- 4. (Original) The compound according to claim 1, wherein R^1 , R^2 and R^3 are each independently-SO₃H or-PO₃H₂.
- 5. (Cancelled)
- 6. (Cancelled)
- 7. (Cancelled)
- 8. (Cancelled)
- 9. (Cancelled)
- 10. (Canceled)
- 11. (Previously Presented) The compound wherein the compound is 4'- (N,N-dimethylamino)-5, 6,7-trimethoxyflavone, 4'- (methylamino)-5, 6,7-trimethoxyflavone, 4'- [N-methyl-N-(3-methoxypropyl)amino)-5,6,7-trimethoxyflavone, 4'- [N,N-di-(2-hydroxyethyl)-amino)-5,7-dihydroxy-6-methoxyflavone, 4'- (2-hydroxyethylamino)-5,7-dihydroxy-6-methoxyflavone, 4'- [2-(N,N-diethylamino)ethylamino]-5,7-dihydroxy-6-methoxyflavone, 4'- [2-(N,N-diethylamino)ethylamino]-5,7-dihydroxy-6-methoxyflavone, 2,3-diphenyl-5,6,7-trimethoxychromone, 2,3-diphenyl-5,6,7-trihydroxychromone, 4'- (methylsulfonamido)-5,6,7-trimethoxyflavone, 4'- [2-(N,N-diethylamino)ethoxy]-6,7-methylenedioxy-5-hydroxy-flavone, 4'- (2,3-dihydroxy-propyloxy)-5,6,7-trimethoxyflavone, or 4'- (Carbmethoxymethoxy)-5,6,7-trimethoxyflavone.
- 12. (Original) A pharmaceutical formulation comprising a compound according to claim 1 and at least one pharmaceutically acceptable carrier, diluent, or excipient.
- 13. (Cancelled)

	ying progression of a condition associated with overproduction of TNF-a selected fror group consisting of rheumatoid arthritis, Crohn's disease, and ulcerative colitis,
com	prising administering to a subject in need thereof an effective amount of a compound ording to claim 1.
15.	(Cancelled)
16.	(Cancelled)
17,	(Cancelled)
18.	(currently amended) A method of treating improving conditions in or delaying
proj	ression of a condition liver damage, lung damage or kidney damage or combinations
ther	eof resulting from over production of TNF-α or superoxide anion raidacals comprising
adm	inistering to a subject in need thereof an effective amount of a compound according to
clai	n l.
19.	(Cancelled)
	(Cancelled) (Cancelled)
21.	
21. 22.	(Cancelled)
21.22.23.	(Cancelled) (Cancelled)

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- 26. (Cancelled)
- 27. (Cancelled)
- 28. (Cancelled)
- 29. (Cancelled)
- 30. (Cancelled)
- 31. (currently amended) A method of treating conditions improving conditions in or delaying progression of a condition selected from liver damage, lung damage or kidney damage the group consisting of diseases associated with the overproduction of TNF-a, everproduction of superoxide anion radical and combinations thereof, comprising administering to a subject in need thereof, a pharmaceutical composition comprising a therapeutically effective amount of a compound of the formula V:

wherein: R^7 , R^8 , and R^9 -are each independently H, lower alkyl,- SO_3H ,- PO_3H_2 ; or R^7 and R^8 together may form a 5-7-membered heterocycle or

X¹ is a substituent on the ortho, meta, or para position of the phenyl ring and is H, C, NH₂, NHCOCH₃, or OR¹⁰, wherein R¹⁰ is H, or lower alkyl, and X¹ is attached in any one of the ortho, meta and para positions or pharmaceutically acceptable salts thereof.

- 32. (Cancelled)
- 33. (Cancelled)
- 34. (Cancelled)
- 35. (Cancelled)
- 36. (Cancelled)

- 37. (Cancelled)
- 38, (Cancelled)
- 39. (previously presented) The method according to claim 31, wherein the compound is 5,6,7- trihydroxyisoflavone, 4',5,6,7- tetrahydroxyflavone, or 4'-amino -5,7-dihydroxy-6-methoxy flavone.
- 40. (Cancelled)
- 41. (Cancelled)
- 42. (Cancelled)
- 43. (Cancelled)
- 44. (previously presented) The method according to claim 31, wherein the pharmaceutical composition is administered in combination with at least one other therapeutic agent useful for the treatment of conditions associated with overproduction of TNF- α , and liver damage, lung damage or kidney damage.
- 45. (Original) The method according to claim 31, wherein the pharmaceutical composition is administered orally or parenterally.
- 46. (currently amended) A method of treating- improving conditions in or delaying progression of a condition conditions selected from the group consisting of diseases associated with the overproduction of TNF-α and combinations thereof, comprising administering to a subject in need thereof, a pharmaceutical composition comprising a therapeutically effective amount of a compound selected from the group consisting of baicalein-6-sulfate, baicalein-6,7-disulfate, bacalein-6-phosphate, bacalein-6,7-diphosphate, baicalein-5,6, 7-triphosphate, sodium and potassium salt derivatives thereof, and pharmaceutically acceptable salts thereof.
- 47. (Cancelled)
- 48. (Cancelled)
- 49 (Cancelled)

- 50 (Cancelled)
- 51. (Cancelled)
- 52. (previously presented) The method according to claim 46, wherein the pharmaceutical composition is administered in combination with at least one other therapeutic agent useful for the treatment of conditions associated with overproduction of TNF- α .
- 53. (Original) The method according to claim 44, wherein the pharmaceutical composition is administered orally or parentally.
- 54. (currently amended) A method of treating conditions improving conditions in or delaying progression of a condition selected from the group consisting of diseases associated with the overproduction of TNF-α, and combinations thereof, comprising administering to a subject in need thereof, a pharmaceutical composition comprising a therapeutically effective amount of compound as in Claim 11.
- 55. (Cancelled)
- 56. (Cancelled)
- 57. (Cancelled)
- 58. (Cancelled)
- 59. (Cancelled)
- 60. (currently amended) A method of treating- improving conditions in or delaying progression of liver damage, lung damage or kidney damage resulting from over production of TNF-α or superoxide anion raidacals which comprises administering to a subject in need thereof a therapeutically effective amount of a compound of the formula:

wherein \mathbb{R}_{4} \mathbb{R}^{1} is selected from hydrogen and alkyl;

R² is selected from hydrogen, lower alkyl and sulfate;

 $\underline{\mathbf{R}^{3}}$ is selected from hydrogen, lower alkyl and sulfate;

 \underline{X}^1 is selected from hydrogen, phenyl and substituted phenyl wherein the substituent is hydroxyl, alkoxy, amino, mono or dialkyl substituted amino, hydroxyl alkoxy, or aminoalkoxy

and X² is selected from hydrogen and phenyl, X¹ and X² cannot both be phenyl.

61 (new) A pharmaceutical formulation comprising a compound according to claim 11 and at least one pharmaceutically acceptable carrier, diluent, or excipient.